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HMRhoNox-M, Fe(II)-selective fluorescent probe

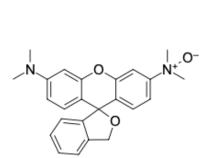
http://www.lumiprobe.com/p/hmrhonox-m

HMRhoNox-M (also known as LysoRhoNox) is a Fe^{2+} -selective fluorescent probe based on the N-oxide-controlled spirocyclization of tetramethyl-hydroxymethyl rhodamine.

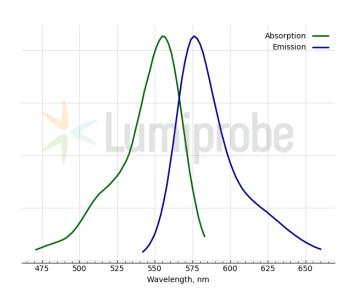
In the absence of Fe^{2+} , HMRhoNox-M exists in the non-fluorescent spirocyclic form showing only negligible fluorescence in an aqueous buffer and at physiological pH. The addition of Fe^{2+} induces a 60-fold increase of the fluorescence signal at 575 nm through the deoxygenation of the dialkylamino group and the transition of the probe to an open fluorescent form. HMRhoNox-M responds to Fe^{2+} in a dose-dependent manner.

The fluorescence response of HMRhoNox-M is highly selective for Fe^{2+} over other transition metal ions, including Fe^{3+} , alkali metal ions, and alkaline earth metal ions.

HMRhoNox-M is the cell-permeant probe that is mainly localized in lysosomes. It is suitable for monitoring fluctuations of endogenous labile iron in living cells, including the transferrin-induced Fe uptake.



Structure of HMRhoNox-M



Absorption and emission spectra of HMRhoNox-M

General properties

Appearance: beige-pinkish crystals

 $\label{eq:molecular weight: 388.47} \mbox{Molecular formula:} \mbox{C_{24}H}_{24}\mbox{N_2O}_3$

Quality control: NMR ¹H and HPLC-MS (95+%)

Storage conditions: 24 months after receival at -20°C in the dark. Transportation: at room temperature

for up to 3 weeks. Desiccate.

Spectral properties

Excitation/absorption maximum, nm: 555
Emission maximum, nm: 575